[DISCUSSION DRAFT]

March 29, 2011

112тн (CONGRESS
1st	Session

H.R.

To amend the consumer product safety laws... $\llbracket to\ be\ provided
rbracket$

IN THE HOUSE OF REPRESENTATIVES

	introduced	the	following	bill;	which	was	referred	to
the Committee	on							

A BILL

To amend the consumer product safety laws... [to be provided]

Be it enacted by the Senate and House of Representatives of the United States of America in Congress assembled,

SECTION 1. DEFINITION OF CHILDREN'S PRODUCT.

(a) DEFINITION.—Section 3(a)(2) of the Consumer

Product Safety Act (15 U.S.C. 2052(a)(2)) is amended—

(1) in the matter preceding subparagraph (A)—

(A) by striking "intended primarily for children 12 years of age or younger" and in-

1	serting "primarily intended for use by children
2	[] years of age or younger'; and
3	(B) by striking "intended for a child 12
4	years of age or younger" and inserting "in-
5	tended for use by a child [] years of age
6	or younger";
7	(2) in subparagraph (B), by striking "children
8	12 years of age or younger" and inserting "children
9	[] years of age or younger"; and
10	(3) in subparagraph (C), by striking "child 12
11	years of age or younger" and inserting "child
12	[] years of age or younger".
13	(b) Technical Amendment.—Section 101(a)(1) of
14	the Consumer Product Safety Improvement Act of 2008
15	(15 U.S.C. 1278a(a)(1)) is amended by striking "(as de-
16	fined in section 3(a)(16) of the Consumer Product Safety
17	Act (15 U.S.C. 2052(a)(16)))" and inserting "(as defined
18	in section 3(a) of the Consumer Product Safety Act (15
19	U.S.C. 2052(a)))".
20	SEC. 2. APPLICATION OF LEAD LIMIT.
21	(a) Extension of Deadline for Lead Limit.—
22	Section 101(a)(2) of the Consumer Product Safety Im-
23	provement Act of 2008 (15 U.S.C. 1278a(a)(2)) is amend-
24	ed

1	(1) in subparagraph (C), by striking "3 years"
2	and inserting "4 years"; and
3	(2) in subparagraph (D), by striking "3 years"
4	and inserting "4 years".
5	(b) Authority to Apply Limits to Other Prod-
6	UCTS.—Such section is further amended by adding at the
7	end the following:
8	"(F) AUTHORITY TO APPLY LIMITS TO
9	OTHER PRODUCTS.—The Commission may, by
10	regulation, apply the limit set forth in subpara-
11	graph (A) to any consumer product other than
12	a children's product (as such terms are defined
13	in section 3(a) of the Consumer Product Safety
14	Act (15 U.S.C. 2052(a))) that is designed or
15	primarily intended for use by children 12 years
16	of age or younger, or to any class of such con-
17	sumer products, if it determines after a hearing
18	that the lead content in such product or class
19	of products presents an unreasonable risk to
20	children's health.".
21	(c) Prospective Application of Lead Limit for
22	CHILDREN'S PRODUCTS.—Section 101(a) of the Con-
23	sumer Product Safety Improvement Act of 2008 (15
24	U.S.C. 1278a(a)) is further amended by adding at the end
25	the following:

1	"(3) APPLICATION.—Each limit set forth in
2	paragraph (2) shall apply only to a children's prod-
3	uct (as defined in section 3(a) of the Consumer
4	Product Safety Act (15 U.S.C. 2052(a))) that is
5	manufactured after the effective date of such respec-
6	tive limit.".
7	(d) Alternative Limit and De Minimis Excep-
8	TION.—Section 101(b) of such Act (15 U.S.C.
9	1278a(b)(1)) is amended—
10	(1) by redesignating paragraphs (2) through 5
11	as paragraphs (3) through (6), respectively; and
12	(2) by striking paragraph (1) and inserting the
13	following:
14	"(1) Alternative limit for certain mate-
15	RIALS AND DE MINIMIS EXCEPTION FOR CERTAIN
16	PARTS.—
17	"(A) Alternative limit.—For a compo-
18	nent part of a children's product that is made
19	of steel, copper, or aluminum alloys, the limit
20	referred to in subsection (a)(1) shall be $\llbracket ___ \rrbracket$
21	parts per million unless—
22	"(i) the product into which such part
23	is incorporated fits entirely within the
24	small parts cylinder described in section

1	1501.4 of title 16, Code of Federal Regula-
2	tions; or
3	"(ii) after any necessary assembly of
4	the product and after the product has been
5	subjected to reasonably foreseeable condi-
6	tions of use and abuse, the part or any
7	portion of the part becomes detached from
8	the product and such part or portion of the
9	part fits entirely within such cylinder.
10	"(B) DE MINIMIS EXCEPTION.—
11	"(i) In general.—The limits estab-
12	lished under subsection (a) shall not apply
13	to any component part of a children's
14	product if, under reasonably foreseeable
15	conditions of use and abuse, it is unlikely
16	that a child who is exposed to the product
17	would ingest more than a de minimis
18	amount of lead, unless—
19	"(I) the product into which such
20	part is incorporated fits entirely with-
21	in the small parts cylinder described
22	in section 1501.4 of title 16, Code of
23	Federal Regulations; or
24	"(II) after any necessary assem-
25	bly of the product and after the prod-

1	uct has been subjected to reasonably
2	foreseeable conditions of use and
3	abuse, the part or any portion of the
4	part becomes detached from the prod-
5	uct and such part or portion of the
6	part fits entirely within such cylinder.
7	"(ii) Methodology for estimating
8	AMOUNT OF LEAD INGESTED.—
9	"(I) Establishment by com-
10	MISSION.—The Commission shall, by
11	regulation, establish a methodology
12	for estimating the amount of lead a
13	child would likely ingest from expo-
14	sure to a component part. Such meth-
15	odology shall distinguish, at a min-
16	imum, between parts that can be
17	placed in the mouth and parts that
18	cannot be placed in the mouth.
19	"(II) Interim methodology.—
20	Until the Commission has issued a
21	final rule under subclause (I), a man-
22	ufacturer may use any reasonable
23	methodology to estimate the amount
24	of lead a child would likely ingest
25	from exposure to a component part.

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1	The manufacturer shall document the
2	methodology used.
3	"(iii) DE MINIMIS AMOUNT DE-
4	FINED.—In this subparagraph, the term
5	'de minimis amount' means []
6	micrograms per day. The Commission may
7	revise such amount by regulation.
8	"(2) Exclusion of Certain used Chil-
9	DREN'S PRODUCTS.—
10	"(A) GENERAL EXCLUSION.—The lead
11	limits established under subsection (a) shall not
12	apply to a used children's product.
13	"(B) Definition.—The term 'used chil-
14	dren's product' means a children's product that
15	was obtained by the seller for use and not for
16	the purpose of resale or was obtained by the
17	seller, either directly or indirectly, from a per-
18	son who obtained such children's product for
19	use and not for the purpose of resale. Such
20	term also includes a children's product that was
21	donated to the seller for charitable distribution
22	or resale to support charitable purposes. Such
23	term shall not include—
24	"(i) children's metal jewelry; or

1	"(ii) any children's product for which
2	the donating party or the seller has actual
3	knowledge that the product is in violation
4	of the lead limits in this section; or
5	For purposes of this definition, the term 'seller'
6	includes a person who lends or donates a used
7	children's product.".
8	SEC. 3. APPLICATION OF THIRD PARTY TESTING REQUIRE-
9	MENTS.
10	(a) Applicable Children's Products.—Section
11	14(a) of the Consumer Product Safety Act (15 U.S.C.
12	2063(a)) is amended—
13	(1) in paragraph (2)—
14	(A) in the matter preceding subparagraph
15	(A), by inserting "described in clauses (i)
16	through (iv) of paragraph (3)(B)" after "a chil-
17	dren's product safety rule';
18	(B) in subparagraph (B), by striking "the
19	children's product safety rule" and inserting
20	"such children's product safety rule"; and
21	(C) by striking the flush sentence following
22	subparagraph (B); and
23	(2) in paragraph (3)—
24	(A) in subparagraph (A), by inserting "de-
25	scribed in clauses (i) through (iv) of subpara-

1	graph (B)" after "a children's product safety
2	rule"; and
3	(B) in subparagraph (B), by striking
4	clauses (v) and (vi).
5	(b) Third Party Testing Requirements.—Sec-
6	tion 14(b) of the Consumer Product Safety Act (15 U.S.C.
7	2063(b)) is amended to read as follows:
8	"(b) Testing Programs.—
9	"(1) In general.—The Commission may, by
10	rule, prescribe reasonable testing programs to be
11	used as the basis for certification under subsection
12	(a).
13	"(2) Testing by an independent third
14	PARTY.—Any test or testing program on the basis of
15	which a certificate is issued under subsection (a)
16	may, at the option of the person required to certify
17	the product, be conducted by an independent third
18	party qualified to perform such tests, unless the
19	Commission, by rule and in accordance with para-
20	graph (3), requires testing by an independent third
21	party for—
22	"(A) a particular rule, regulation, stand-
23	ard, ban;
24	"(B) any portion of a particular rule, regu-
25	lation, standard, or ban; or

1	"(C) a particular class of products.
2	"(3) Requirements for testing by an
3	INDEPENDENT THIRD PARTY.—The Commission
4	may not require testing by an independent third
5	party under paragraph (2) until the Commission has
6	completed each of the following:
7	"(A) Accreditation of conformity as-
8	SESSMENT BODIES.—Established and published
9	notice of the requirements for accreditation of
10	third party conformity assessment bodies who
11	are determined to be qualified by the Commis-
12	sion to conduct such testing.
13	"(B) Testing Capacity.—Determined
14	that the testing capacity of the accredited third
15	part conformity assessment bodies is sufficient
16	to prevent unreasonable delays due to testing.
17	"(C) Exemptions and alternate test-
18	ING PROCEDURES.—
19	"(i) In general.—Established, by
20	rule—
21	"(I) exemptions for works of art
22	and other one-of-a-kind products; and
23	"(II) exemptions or alternative
24	testing procedures for the certification
25	of specialty products for the disabled,

1	and products that are produced in
2	small quantities such that the cost of
3	testing by an independent third party
4	is not economically practicable.
5	"(ii) Produced in small quan-
6	TITIES DEFINED.—In this subparagraph,
7	the term 'produced in small quantities'
8	means that not more than [] units of
9	the same product (or substantially similar
10	products) are produced in one year by a
11	manufacturer and any affiliated manufac-
12	turer. A manufacturer may not subdivide
13	the production of such manufacturer into
14	small quantities in order to evade third
15	party testing requirements.
16	"(D) Rulemaking considerations.—
17	Made a reasoned determination—
18	"(i) that the benefits from requiring
19	third-party testing justify the costs (recog-
20	nizing that some costs are difficult to
21	quantify); and
22	"(ii) that any rule issued pursuant to
23	this paragraph is tailored to impose the
24	least possible burden, taking into account

1	to the extent practicable, the costs of cu-
2	mulative regulations.
3	"(4) Review of Previous Rules.—The Com-
4	mission may not enforce a third party testing re-
5	quirement that became effective during the period
6	after August 14, 2009, and before the date of the
7	enactment of the $[$ Act of 2011 $]$
8	(or that was stayed by the Commission during such
9	period) until the Commission has reviewed such re-
10	quirement and promulgated any revisions as nec-
11	essary to ensure compliance with the requirements
12	of paragraph (3).".
13	(c) Continuing Testing.—Section 14(d)(2) of the
14	Consumer Product Safety Act (15 U.S.C. $2063(d)(2)$) is
15	amended—
16	(1) in the matter preceding subparagraph (A),
17	by striking "Not later than 15 months after the date
18	of enactment of the Consumer Product Safety Im-
19	provement Act of 2008, the" and inserting "The";
20	(2) in the matter preceding subparagraph (A),
21	by striking "shall";
22	(3) in subparagraph (A), by striking "initiate"
23	and inserting "not later than 15 months after the
24	date of enactment of the Consumer Product Safety
25	Improvement Act of 2008, shall initiate"; and

1	(4) in subparagraph (B), by striking "estab-
2	lish" and inserting "may establish".
3	SEC. 4. APPLICATION OF AND PROCESS FOR UPDATING DU-
4	RABLE NURSERY PRODUCTS STANDARDS.
5	(a) Application of Standard.—Section 104 of the
6	Consumer Product Safety Improvement Act of 2008 (15
7	U.S.C. 2056a) is amended—
8	(1) in subsection (c), by redesignating para-
9	graph (3) as paragraph (4) and inserting after para-
10	graph (2) the following:
11	"(3) Application.—
12	"(A) In General.—Paragraph (1) shall
13	not apply to any revision of the standard pro-
14	mulgated under subsection (b)(1)(B) subse-
15	quent to the initial promulgation of a standard
16	under such subsection.
17	"(B) Special rule for fixed-side
18	CRIBS SUBJECT TO CERTAIN STATE OR LOCAL
19	LAW REQUIREMENTS.—Paragraph (1) shall not
20	apply to a fixed-side crib offered or provided for
21	use in a licensed child care facility that is sub-
22	ject to the following requirements under the law
23	of a State or a political subdivision of a State:

1	"(i) The facility may not allow a child
2	to remain in a crib for any significant
3	amount of time while the child is awake.
4	"(ii) The facility may not place in a
5	crib a child over the age of [12 months].
6	"(iii) An adult must be present when-
7	ever a child is in a crib.".
8	(b) UPDATING STANDARD.—Section 104(b) of the
9	Consumer Product Safety Improvement Act of 2008 (15
10	U.S.C. 2056a(b)) is amended by adding at the end the
11	following:
12	"(4) Process for considering subsequent
13	REVISIONS TO VOLUNTARY STANDARD.—
14	"(A) NOTICE OF ADOPTION OF VOL-
15	UNTARY STANDARD.—When the Commission
16	promulgates a consumer product safety stand-
17	ard under this subsection that is based, in
18	whole or in part, on a voluntary standard, the
19	Commission shall notify the organization that
20	issued the voluntary standard of the Commis-
21	sion's action and shall provide a copy of the
22	consumer product safety standard to the orga-
23	nization.
24	"(B) Commission action on revised
25	VOLUNTARY STANDARD.—If an organization re-

1	vises a standard that has been adopted, in
2	whole or in part, as a consumer product safety
3	standard under subparagraph (A), it shall no-
4	tify the Commission. The revised voluntary
5	standard shall be considered to be a consumer
6	product safety standard issued by the Commis-
7	sion under section 9 of the Consumer Product
8	Safety Act (15 U.S.C. 2058), effective 180 days
9	after the date on which the organization notifies
10	the Commission (or such later date specified by
11	the Commission in the Federal Register) unless,
12	within 90 days after receiving that notice, the
13	Commission notifies the organization that it has
14	determined that the proposed revision does not
15	improve the safety of the consumer product cov-
16	ered by the standard and that the Commission
17	is retaining the existing consumer product safe-
18	ty standard.".
19	SEC. 5. APPLICATION OF SECTION 106 TO FDA-REGULATED
20	PRODUCTS.
21	Section 106(a) of the Consumer Product Safety Im-
22	provement Act (15 U.S.C. 2056b(a)) is amended by in-
23	serting "or any provision that restates or incorporates a
24	regulation promulgated by the Food and Drug Adminis-

1	tration or any statute administered by the Food and Drug
2	Administration" after "or by statute".
3	SEC. 6. APPLICATION OF PHTHALATES STANDARD.
4	(a) Prospective Application, Accessible, Plas-
5	TICIZED COMPONENT PARTS.—Section 108 of the Con-
6	sumer Product Safety Improvement Act of 2008 (15
7	U.S.C. 2057c) is amended—
8	(1) by redesignating subsections (c) through (e)
9	as subsections (d) through (f), respectively; and
10	(2) by inserting after subsection (b) the fol-
11	lowing:
12	"(c) Application.—
13	"(1) Accessible component parts.—Sub-
14	sections (a) and (b)(1) and any rule promulgated
15	under subsection (b)(3) shall apply to any children's
16	toy or child care article containing any accessible,
17	plasticized component part that is manufactured
18	after the respective effective dates in each such sub-
19	section and any such final rule.
20	"(2) Commission authority.—The Commis-
21	sion may, by rule, exempt any children's toy or child
22	care article described in paragraph (1) or any class
23	of such products or materials used in such products
24	from any of the prohibitions under subsections (a)
25	and (b)(1) and any rule promulgated under sub-

1	section (b)(3) where the Commission determines that
2	compliance with any such prohibition is not nec-
3	essary to protect children's health.".
4	(b) Effect of Conclusions of the Chronic
5	HAZARD ADVISORY PANEL.—Section 108(b)(3) of such
6	Act (15 U.S.C. 2057c(b)(3)) is amended—
7	(1) by striking "Not later than" and inserting
8	the following:
9	"(A) Rulemaking required.—Not later
10	than";
11	(2) by redesignating subparagraphs (A) and
12	(B) as clauses (i) and (ii), respectively;
13	(3) in clause (i) (as so redesignated), by insert-
14	ing "or terminate such prohibition" after "margin of
15	safety"; and
16	(4) by adding at the end the following:
17	"(B) Deadline and effect on prohibi-
18	TION.—If the Commission does not commence a
19	rulemaking proceeding within 90 days after re-
20	ceiving the report required by paragraph (2)(C)
21	or does not issue a final rule as required by
22	subparagraph (A) within [] after
23	receiving such report, the prohibition in para-
24	graph (1) shall terminate.".

1	(c) Definitions.—Section 108(f) of the Consumer
2	Product Safety Improvement Act of 2008 (15 U.S.C.
3	2057c(f)) (as redesignated by subsection (a)) is amend-
4	ed—
5	(1) in paragraph (1)—
6	(A) in subparagraph (B), by striking "con-
7	sumer product" and all that follows and insert-
8	ing "children's product that is subject to the
9	standard made mandatory by section 106(b) or
10	any successor standard";
11	(B) in subparagraphs (C), by striking
12	"consumer product" and inserting "children's
13	product"; and
14	(C) in subparagraph (D)—
15	(i) by striking "consumer product"
16	and inserting "children's product";
17	(ii) by striking "section 3(a)(1)" and
18	inserting "section 3(a)"; and
19	(iii) by striking "2052(a)(1)" and in-
20	serting "2052(a)"; and
21	(2) by amending paragraph (2) to read as fol-
22	lows:
23	"(2) Determination guidelines.—For pur-
24	poses of this section, a toy can be placed in a child's
25	mouth if any part of the toy can actually be brought

1	to the mouth and kept in the mouth by a child so
2	that it can be sucked and chewed. If the children's
3	product can only be licked, it is not regarded as able
4	to be placed in the mouth. If a toy or part of a toy
5	in one dimension is smaller than 5 centimeters, it
6	can be placed in the mouth.".
7	SEC. 7. EXEMPTION AUTHORITY FOR TRACKING LABELS
8	REQUIREMENT.
9	Section 14(a)(5) of the Consumer Product Safety Act
10	(15 U.S.C. 2063(a)(5)) is amended—
11	(1) by striking "Effective 1 year" and inserting
12	"(A) Effective 1 year";
13	(2) by redesignating subparagraphs (A) and
14	(B) as clauses (i) and (ii), respectively; and
15	(3) by adding at the end the following:
16	"(B) The Commission may, by regulation, exclude a
17	specific product or class of products from the require-
18	ments in subparagraph (A) if the Commission determines
19	that it is not economically practicable for such product or
20	class of products to bear the marks required by such sub-
21	paragraph. The Commission may establish alternative re-
22	quirements for any product or class of products excluded
23	under the preceding sentence consistent with the purposes
24	described in clauses (i) and (ii) of subparagraph (A).".

	- v
1	SEC. 8. REQUIREMENTS FOR PUBLIC DATABASE.
2	(a) Requirements for Submissions to the
3	Database.—Section 6A(b) of the Consumer Product
4	Safety Act (15 U.S.C. 2055a(b)) is amended—
5	(1) in paragraph (1)(A)—
6	(A) in clause (i), by striking "consumers"
7	and inserting "persons who suffer harm or risk
8	of harm related to the use of a product, their
9	next of kin or members of their household, their
10	legal representative, or another person expressly
11	authorized by any such person"; and
12	(B) in clause (v), by striking "public safety
13	entities" and inserting "police, fire, ambulance
14	emergency medical services, Federal, State, and
15	local law enforcement entities, and other related
16	public safety officials"; and
17	(2) in paragraph (2)(B)—
18	(A) in clause (i), by inserting "and its lo-
19	cation and availability" after "concerned";
20	(B) in clause (iv), by inserting "and it
21	such person is not the person harmed by the

product, the name and contact information of

the person who suffered the harm or risk of

harm related to the use of the product" after

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"report"; and

1	(C) in clause (v), by inserting "that such
2	person is the consumer who used the product
3	that gave rise to the harm, the user's next of
4	kin, a member of the user's household, the legal
5	representative of the user, or another person ex-
6	pressly authorized by any such person and"
7	after "person submitting the information".
8	(b) ADEQUACY AND ACCURACY OF INFORMATION RE-
9	PORTED TO THE PUBLIC DATABASE.—Section 6A(c)(2) of
10	the Consumer Product Safety Act (15 U.S.C.
11	2055a(c)(2)) is amended—
12	(1) in subparagraph (A), by striking "to sub-
13	mit" and all that follows and inserting "to—
14	"(i) notify the Commission within
15	[] days after receipt of the report
16	that the information provided in the report
17	is insufficient for determining which of the
18	manufacturer's products is the subject of
19	the complaint, in which case the manufac-
20	turer shall provide the Commission (and
21	the person submitting the complaint, if
22	that person has consented to disclosure of
23	contact information) with information to
24	assist the person submitting the report to

1	sufficiently identify or provide an adequate
2	description of the product;
3	"(ii) notify the Commission within
4	[] days after receipt of the report
5	that the information provided in the report
6	is materially inaccurate and to provide the
7	Commission with any additional informa-
8	tion supporting the manufacturer's claim
9	of inaccuracy; and
10	"(iii) submit other comments to the
11	Commission on the information contained
12	in such report."; and
13	(2) by redesignating subparagraphs (B) and
14	(C) as subparagraphs (C) and (D), respectively, and
15	inserting after subparagraph (A) the following:
16	"(B) ACTION BY THE COMMISSION.—
17	"(i) Insufficient information.—If
18	a manufacturer notifies the Commission of
19	the insufficiency of the information in a re-
20	port pursuant to subparagraph (A)(i), the
21	Commission shall provide the information
22	provided by the manufacturer to the per-
23	son submitting the report (unless such in-
24	formation has already been provided di-
25	rectly by the manufacturer) and seek to

1	obtain from such person an adequate de-
2	scription of the product.
3	"(ii) Materially inaccurate in-
4	FORMATION.—If a manufacturer notifies
5	the Commission of a material inaccuracy in
6	a report pursuant to subparagraph (A)(ii),
7	and the Commission determines that the
8	claim is potentially valid, the Commission
9	shall seek to resolve the inaccuracy by any
10	of the following:
11	"(I) Obtaining from the person
12	submitting the report such additional
13	information necessary to correct the
14	inaccuracy.
15	"(II) Investigating the incident
16	giving rise to the report in order to
17	correct any such inaccuracy.
18	"(III) Providing the manufac-
19	turer a reasonable period of time to
20	investigate and provide additional in-
21	formation to correct any inaccuracy.
22	"(iii) Stay on inclusion in data-
23	BASE.—The Commission shall not include
24	in the database a report described in
25	clauses (i) or (ii) until the product can be

1	specifically identified and any material in-
2	accuracy corrected.".
3	(c) Misrepresentation Prohibited.—Section
4	19(a)(13) of the Consumer Product Safety Act by insert-
5	ing "related to a submission of information to the data-
6	base established under section 6A, or" after "misrepresen-
7	tation to such an officer or employee".
8	SEC. 9. SUBPOENA AUTHORITY.
9	Section 27(b) of the Consumer Product Safety Act
10	(15 U.S.C. 2076(b)) is amended—
11	(1) in paragraph (3), by inserting "and phys-
12	ical" after "documentary";
13	(2) in paragraph (8), by striking "and";
14	(3) by redesignating paragraph (9) as para-
15	graph (10) and inserting after paragraph (8) the fol-
16	lowing:
17	"(9) to delegate to the general counsel of the
18	Commission the authority to issue subpoenas solely
19	to Federal, State, or local government agencies for
20	evidence described in paragraph (3); and"; and
21	(4) in paragraph (10) (as so redesignated), by
22	inserting "(except as provided in paragraph (9))"
23	after "paragraph (3)".

I	SEC. 10. AVAILABILITY OF CERTAIN PERSONAL AND MED-
2	ICAL INFORMATION TO THE CPSC.
3	Section 5 of the Consumer Product Safety Act (15
4	U.S.C. 2054) is amended by adding at the end the fol-
5	lowing new subsection:
6	"(e) Availability of Personal and Medical In-
7	FORMATION UNDER HIPAA.—In order to carry out its
8	investigative and enforcement activities under this Act and
9	under any of the Acts enforced by the Commission, the
10	Commission shall be deemed a public health authority
11	within the meaning of section 164.512(b)(i) of title 45,
12	Code of Federal Regulations, for purposes of permitted
13	disclosures of protected health information authorized
14	under such section. For purposes of such section informa-
15	tion about deaths, injuries, diseases, and other health im-
16	pairments possibly relating to consumer products shall be
17	deemed protected health information authorized to be dis-
18	closed to such public health authorities under such sec-
19	tion.".
20	SEC. 11. EFFECTIVE DATE.
21	The amendments made by this Act shall be treated
22	as having taken effect on the date of enactment of the
23	Consumer Product Safety Improvement Act of 2008.